IN THE CLAIMS:

Please enter any changes in the claims indicated in the complete copy of the pending claims, as sought to be amended, presented below:

1. (Currently Amended) A compound of Formula 1:

$$R1 \xrightarrow{H} \begin{array}{c} R2 \\ N \\ O \\ \end{array} \begin{array}{c} R3 \\ N \\ H \\ \end{array} \begin{array}{c} X \\ N \\ Ar_1 \\ \end{array}$$

Formula 1

wherein:

R₁ is selected from cycloalkyl and, heterocycloalkyl, aryl and heteroaryl,

wherein R₁ is optionally substituted with one or more substituents R_a,

wherein R_a is independently selected from the group consisting of alkyl, halo, haloalkyl, nitro, alkenyl, alkynyl, alkoxy, $-(R_7)_nNR_8R_9$ (wherein R_7 is selected from alkyl alkylene, alkoxy alkylene oxide, and oxyalkyl oxyalkylene, R_8 and R_9 can be independently selected from H, and alkyl, or R_8 and R_9 can join together such that NR_8R_9 form a 5 or 6-member heterocyclic ring, and n is selected from 0 and 1), and the substituent R_a is optionally further substituted with one or more substituents selected from the group consisting of alkyl, alkoxy, halo, cyano, alkanoyl, haloalkyl, thioalkyl and nitro, $-(R_7)_nNR_8R_9$, wherein R_7 , R_8 , R_9 , and n are as defined above—;

R₂ and R₃ are: a) independently selected from the group consisting of H, alkyl, haloalkyl, aralkyl optionally substituted aryl, optionally substituted heteroaryl and optionally substituted, saturated or unsaturated, 5-or 6-membered, homocyclic or heterocyclic rings wherein the optional substituent may be selected from the group consisting of H, alkyl, alkoxy, and halo;

Of

b) join together to form a 3, 4, 5, 6 or 7 member spirocyclic ring;

X is selected slelected from O, S, NH and NCN;

 Ar_1 is phenyl and is optionally substituted with one or more substituents R_{b} ,

wherein the substituent(s) R_b are independently selected from the group consisting of alkyl, alkoxy, alkanoyl, nitro halo, haloalkoxy, $-(R_7)_nNR_8R_9$, $-S(O)_2NR_{10}R_{11}$ and $-O-(CH_2)_mNR_{10}R_{11}$ (wherein R_7 is selected from alkyl, alkoxy, and oxyalkyl, R_8 and R_9 can be independently selected from H, and alkyl, or R_8 and R_9 can join together such that NR_8R_9 form a 5 or 6-member heterocyclic ring, and n is selected from 0, 1, 2, 3, 4 and 5 and R_{10} and R_{11} are independently selected from H, or alkyl, or R_{10} and R_{11} can join together such that $NR_{10}R_{11}$ to form a 5 or 6-member heterocyclic ring and m is selected from 1, 2, 3, 4 and 5) and;

the substituent R_b is optionally further substituted with one or more substituents selected from the group consisting of alkyl, alkoxy, halo, cyano, alkanoyl, haloalkyl, thioalkyl, nitro, $-(R_7)_nNR_8R_9$ wherein R_7 , R_8 , R_9 and n are as described above,

with the proviso that Ar_1 does not have a substituent at the 2-position selected from the following groups, nitro, haloalkyl, eyano, $C(O)R_{12}$, $C(O)OR_{12}$, $C(O)NR_{12}R_{13}$, $S(O)R_{12}$, $S(O)_2R_{12}$, and $S(O)_2NR_{12}R_{13}$ (wherein R_{12} and R_{13} are independently selected from H and alkyl), and,

the second proviso that Ar_1 does not have an alkanoyl substituent at the 4 position, and a salt solvate or hydrate thereof.

2. (Currently Amended) A compound of claim 1 wherein Ar₁ is substituted with one or more substituents, \underline{R}_{b} \underline{R}_{a} , wherein the substituent(s) \underline{R}_{b} \underline{R}_{a} are selected from the group consisting of alkyl, alkoxy, nitro, acetyl, halo, haloalkyl, -S(O)₂NR₁₀R₁₁, -O-(CH₂)_nNR₁₀R₁₁, wherein R₁₀ and R₁₁ are independently selected from H, or alkyl, or R₁₀ and R₁₁ can join together such that NR₁₀R₁₁ form a 5 or 6 member heterocyclic ring.

- 3. (Currently Amended) A compound of claim 2 wherein there are two substituents \underline{R}_b \underline{R}_6 , independently selected from the group consisting of nitro, methoxy, and ethoxy.
- 4. (Currently Amended) A compound of claim 3 wherein the two substituents \underline{R}_b \underline{R}_6 are a nitro substituent at the 5-position and a methoxy substituent at the 2-position.
- (Currently Amended) A compound as defined in claim 1 wherein R₁ is optionally substituted and is selected from the group consisting of phenyl, naphthyl, tetrahydronaphthyl, and indanyl, quinolinyl and pyridyl.
- 6. (Original) A compound of claim 5 wherein R_1 is indanyl.
- 7. (Original) A compound of claim 5 wherein R_1 is optionally substituted pyridyl wherein the substituent(s) R_a are selected from the group consisting of alkyl, and haloalkyl.
- 8. (Original) A compound of claim 5 wherein R₁ is optionally substituted phenyl wherein the substituent(s) R_a are selected from the group consisting of alkyl, halo, haloalkyl, nitro, vinyl, alkoxy, -(R₇)_nNR₈R₉ wherein R₇ is selected from alkyl, alkoxy, and oxyalkyl, R₈ and R₉ can be independently selected from H, and alkyl, or R₈ and R₉ can join together such that NR₈R₉ form a heterocyclic ring, and n is selected from 0 and 1.
- 9. (Original) A compound of claim 8 wherein R₁ is selected from mono or di-substituted phenyl with the substituents selected independently from the group consisting of alkyl, halo and haloalkyl.
- 10. (Currently Amended) A compound as defined in claim 1 wherein R₂ and R₃ are independently selected from, H, alkyl, <u>haloalkyl</u>, aralkyl, optionally substituted aryl, optionally substituted heteroaryl and optionally substituted saturated or unsaturated 5 or 6-membered homocyclic, or heterocyclic rings.
- 11. (Original) A compound as defined in claim 10 wherein R₂ and R₃ are selected independently from H, phenyl, 3-thiophene, sec-butyl, 3,4-difluorophenyl, cyclohexyl, 3-trifuoromethylphenyl, t-butyl, isopropyl, methyl, benzyl, trifuoromethyl.
- 12. (Canceled).
- 13. (Currently Amended) A compound of claim 1 selected from the group consisting of:

- 2-[3-(2-methoxy-5-nitro-phenyl)-thioureido]- N-(2-indanyl)-2-(3-thienyl) acetamide E42.2;
- 2-[3-(2-methoxy-5-nitro-phenyl)-thioureido]- N-(3,4-dimethylphenyl)-2-phenyl acetamide E32.2;
- 2-[3-(2-methoxy-5-nitro-phenyl)-ureido]- N-(3,4-dimethylphenyl)-2-phenyl acetamide E32.5;
- (R)-2-[3-(2-methoxy-5-nitro-phenyl)-thioureido]- N-(3,4-dimethylphenyl)-2-phenyl acetamide E33.1*;
- 2-[3-(2-methoxy-5-nitro-phenyl)-ureido]- N-(2-indanyl)-2-(3-thienyl) acetamide E42.1;
- (R) 2-[3-(2-nitro-5-methoxy-phenyl) ureido] N-(2-indanyl) 2-phenyl R-N-(indan-5-yl)-2-[3-(2-methoxy-5-nitro-phenyl)-ureido]2-phenyl acetamide **E29.1***;
- (R) 2-[3 (2-nitro-5-methoxy-phenyl) ureido]- N (4-chlorophenyl) 2-phenyl R-2-[3-(2-methoxy-5-nitro-phenyl)-ureido]-N-(4-chlorophenyl)-2-phenyl acetamide E4.1; and
- (R)-2-[3-(2-methoxy-5-nitro-phenyl)-ureido]- N-(3-trifluromethylphenyl)-2-phenyl acetamide **E31.2.**
- 14. (Original) A pharmaceutical composition comprising a therapeutically effective amount of a compound of claim 1 and a pharmaceutically acceptable carrier.
- 15. (**Original**) A method for treating a patient having a medical condition for which a glycine transport inhibitor is indicated, comprising the step of administering to a patient a pharmaceutical composition as described in claim 14.
- 16. (Original) A method according to claim 15 wherein the medical condition is schizophrenia, cognitive dysfunction, or Alzheimer's disease.
- 17. (New) A pharmaceutical composition of claim 14 wherein Ar_1 is substituted with one or more substituents, R_b , wherein the substituent(s) R_b are selected from the group consisting of alkyl, alkoxy, nitro, acetyl, halo, haloalkyl, $-S(O)_2NR_{10}R_{11}$, $-O-(CH_2)_nNR_{10}R_{11}$, wherein R_{10} and R_{11} are independently selected from H, or alkyl, or R_{10} and R_{11} can join together such that $NR_{10}R_{11}$ form a 5 or 6 member heterocyclic ring.

- 18. (New) A pharmaceutical composition of claim 14 wherein there are two substituents R_b, independently selected from the group consisting of nitro, methoxy, and ethoxy.
- 19. (New) A pharmaceutical composition of claim 14 wherein the two substituents R_b are a nitro substituent at the 5-position and a methoxy substituent at the 2-position.
- 20. (New) A pharmaceutical composition of claim 14 wherein R₁ is optionally substituted and is selected from the group consisting of phenyl, naphthyl, tetrahydro-naphthyl and indanyl
- 21. (New) A pharmaceutical composition of claim 14 wherein R₂ and R₃ are independently selected from, H, alkyl, haloalkyl, aralkyl, optionally substituted aryl, optionally substituted heteroaryl and optionally substituted saturated or unsaturated 5 or 6-membered homocyclic, or heterocyclic rings.